

Notice of References Cited	Application/Control No.	Applicant(s)/Patent Under Reexamination	
	10/657,762	CRISTALLI, GLORIA	
	Examiner L. E. Crane 	Art Unit 1623	Page 1 of 2

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*	Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
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NON-PATENT DOCUMENTS

*	Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)		
U	(R) Volpini et al., "N6-Alkyl-2-alkynyl Derivatives of Adenosine as Potent and Selective Agonists at the Human Adenosine A3 Receptor and a Starting Point for Searching A2B Ligands," Journal of Medicinal Chemistry, 45(15), 3271-3279 (July 18, 2002)		
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W	(T) Siddiqi et al., "Search for New Purine- and Ribose-Modified Adenosine Analogues as Selective Agonists and Antagonists a Adenosine Receptors," Journal of Medicinal Chemistry, 38(7), 1174-1188 (March 31, 1995).		
X	(U) Klotz et al. (I), "2-Substituted N-ethylcarboxamidoadenosine Derivatives as High-Affinity Agonists at A3 Adenosine Receptors," Naunyn-Schmiedeberg's Archives of Pharmacology, 360(2), 103-108 (1999); published online on July 13, 1999.		

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*	V	(W) Stu Borman, "A3 Receptors," Science & Technology Section of Chemical & Engineering News, 79(7), 37-40 (February 12, 2001).
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Notice of References Cited		APPLICANT(S) Cristalli et al.			

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Foreign Patent Documents

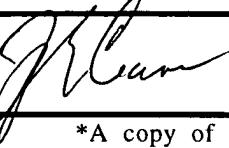
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S	Baraldi et al., "Synthesis and Biological Activity of a New Series of N ⁶ -Arylcarbamoyl, 2(Ar)alkynyl-N ⁶ -arylcaramoyl, and N ⁶ -Carboxamido Derivatives of Adenosine-5'-N-ethyluronamide as A ₁ and A ₃ Adenosine Receptor Agonists," <i>Journal of Medicinal Chemistry</i> , 41(17), 3174-3185 (Aug. 13, 1998).

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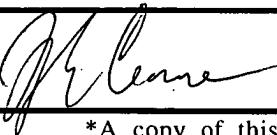
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V†	Klotz et al. (II), "Comparative Pharmacology of Human Adenosine Receptor Subtypes - Characterization of Stably Transfected Receptors in CHO Cells," <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 357(1), 1-9 (1998).
* W	Stu Borman, "A ₃ Receptors," <i>Science & Technology Section of Chemical & Engineering News</i> , 79(7), 37-40 (February 12, 2001).††

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